#9/B
1646
9-28-01

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re the application of:

Tschope, et. al. Atty. Docket No.: 100564-00002

Application Number: 09/508,510 Group Art Unit: 1646

Filed: May 26, 2000 Examiner: Prasad, Sarada

For: LIQUID INTERFERON-ß FORMULATIONS

STATEMENT UNDER 37 C.F.R. § 1.825

Commissioner of Patents
Washington, D.C. 20231

September 24, 2001

Sir:

In accordance with 37 C.F.R. § 1.825, applicant hereby submits the revised Sequence Listing for the above-referenced application in both paper copy and in computer readable form.

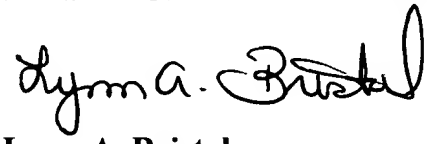
The name of the file on the computer readable form is 100564-00002.txt. The paper copy and the computer readable form are the same. Please note that no new matter has been added.

In the event that this paper is not considered timely filed, applicant hereby petitions for an appropriate extension of time. If necessary, please charge any

additional amounts or credit any overpayments to Direct Deposit Account Number 01-2300.

Respectfully Submitted,

ARENT FOX KINTNER PLOTKIN & KAHN, PLLC



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SEP 14 2001

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UNITED STATES DEPARTMENT OF COMMERCE
Patent and Trademark Offic
Address: COMMISSIONER OF PATENTS AND TRADEMARKS
Washington, D.C. 20231

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.
09/509,810	06/26/00	TSCHOPP	M P100564-0000

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EXAMINER	
PRASAD,S	
ART UNIT	**PAPER NUMBER**
1646	8

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DATE MAILED: 09/13/01

LAB 100564 00002

Oct 13, 2001

Please find below and/or attached an Office communication concerning this application or proceeding.

Commissioner of Patents and Trademarks

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DATE _____

PTO-90C (Rev. 2/95)

2 - Mail Copy

*U.S. GPO: 2000-473-000/44602

Notice to Comply	Application No. 09508510	Applicant(s) Tschope et al
	Examiner Sarada Prasad	Art Unit 1646

NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Applicant must file the items indicated below within the time period set the Office action to which the Notice is attached to avoid abandonment under 35 U.S.C. § 133 (extensions of time may be obtained under the provisions of 37 CFR 1.136(a)).

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

☒ 1. This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to the final rulemaking notice published at 55 FR 18230 (May 1, 1990), and 1114 OG 29 (May 15, 1990). If the effective filing date is on or after July 1, 1998, see the final rulemaking notice published at 63 FR 29620 (June 1, 1998) and 1211 OG 82 (June 23, 1998).

☐ 2. This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c).

☐ 3. A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).

☒ 4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing."

☒ 5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).

☐ 6. The paper copy of the "Sequence Listing" is not the same as the computer readable from of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).

☐ 7. Other:

Applicant Must Provide:

☒ An initial or substitute computer readable form (CRF) copy of the "Sequence Listing".

☒ An initial or substitute paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.

☒ A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).

For questions regarding compliance to these requirements, please contact:

For Rules Interpretation, call (703) 308-4216
For CRF Submission Help, call (703) 308-4212
PatentIn Software Program Support
Technical Assistance.....703-287-0200
To Purchase PatentIn Software.....703-306-2600

PLEASE RETURN A COPY OF THIS NOTICE WITH YOUR REPLY

SEQUENCE LISTING

<110> Tschope, Michael
Siklosi, Thomas
Schroeder, Peter
Hofer, Hans

<120> Liquid Interferon-B Formulations

<130> 100564-00002

<140> 09/508,510
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